

Hydrogen For A PEM Fuel Cell Vehicle Using A Chemical-Hydride Slurry

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INTRODUCTION

Because of the inherent advantages of high efficiency, environmental acceptability, and high modularity, fuel cells are potentially attractive power suppliers. Worldwide concerns over clean environments have revitalized research efforts on developing fuel cell vehicles (FCVs). As a result of intensive research efforts, most of the subsystem technologies for FCVs are currently well established. These include: high power density PEM fuel cells, control systems, thermal management technology, and secondary power sources for hybrid operation. For mobile applications, however, supply of hydrogen or fuel for fuel cell operation poses a significant logistic problem.

A great many technologies have been investigated as candidates for the onboard storage of pure hydrogen for FCVs. These technologies include: 1) compressed hydrogen, 2) liquefied hydrogen, 3) rechargeable metal hydride, 4) carbon adsorption and hybrid systems, and 5) liquid hydrides and other chemical hydrides. However, the volume and/or weight energy densities of these onboard hydrogen storage technologies are significantly lower than those of internal combustion engines or the DOE hydrogen plan. Therefore, development of a high energy density subsystem to supply hydrogen for fuel cell operation is an urgently needed technology for the successful development of FCVs.

To supply high-purity hydrogen for FCV operation, Thermo Power Corporation (THP) has developed an advanced hydrogen storage technology. In this approach, a chemical (light metal) hydride/organic slurry is used as the hydrogen carrier and storage media. At the point of use, high purity hydrogen will be produced by reacting the hydride/organic slurry with water. The fluid-like nature of the hydride/organic slurry will provide us a unique opportunity for pumping, transporting, and storing these materials. In addition, the spent hydride can relatively easily be collected at the pumping station and regenerated utilizing renewable sources, such as biomass, natural gas, or coal at the central processing plants. Therefore, the entire process will be economically favorable and environmentally friendly. The final product of the program is a user-friendly and relatively high energy storage density hydrogen supply system for fuel cell operation.

BACKGROUND AND TECHNICAL APPROACH

Pros and cons of the currently available and advanced hydrogen storage technologies, along with expected performance of the proposed technology, are summarized in Table 1. A plot showing how chemically-reacting hydrides compare with other fuels is shown in Figure 1.

An essential feature of the THP approach is to develop a relatively high energy storage density hydrogen supply system based on exothermic chemical reactions between metal hydrides and water. Hydrogen production via metal hydride and water reactions is a well-established industrial process. In fact, several groups of researchers have investigated the metal hydride/water reaction process to supply hydrogen for fuel cells for mobile power generations. In this research, it has been identified that reaction rate control, frequent on/off operation, and safety of the operation could be significant problems for high energy density operations.

One of the key technical challenges in the program is, therefore, to precisely control the metal hydride and water reaction. In our approach, the continuous organic slurry media will act as a path for dissipating heat that is generated from the hydride/water reaction. Furthermore, by controlling surface chemistry of the organic media, the water/metal hydride reaction rate can easily be controlled. This concept is shown in Figure 2. In Figure 2a, a sketch is shown of two hydride particles, one surrounded by oil and one not. The oil layer inhibits the water access to the hydride and thereby controls the rate of reaction, which would otherwise be explosive. In Figure 2b, the hydride suspension is shown to exemplify how the dispersant acts to hold the particle in suspension within the oil and further inhibit the reaction with water.

Because of the reaction rate control afforded by the organic media, the hydrogen reactor can be a simple device. Water and hydride slurry are metered into the reactor, where they are thoroughly mixed to ensure complete reaction. This reaction goes to completion quickly, leaving a powdery waste. Hydrogen production rate is controlled by the injection rate of water and hydride. Heat

released by the reaction can be absorbed by the evaporation of water. No complicated control systems are needed to ensure proper and safe operation of the hydrogen reactor.

The water required for thermal control and hydrogen reaction is provided by condensed vapor from the hydrogen fuel cell. Only a small reservoir of water is required for startup, makeup, and surge demand. Thus, the required water does not significantly affect the volumetric and gravimetric energy storage densities.

The slurry form of hydride has other benefits beyond reaction control. The hydride fuel can be handled as a liquid, simplifying transportation, storage, and delivery. Use of a slurry permits refueling similar to current gasoline filling stations, allowing the tank to be easily topped-off at any time. The hydroxide waste products produced by the hydrogen system can be washed from the onboard storage tank during the slurry filling operation. Both the hydride fuel and hydroxide waste product can be easily transported between the distribution centers and a central recycling plant.

The used reactant slurry containing LiOH is returned to a central processing plant where the LiOH is recycled to LiH in a large-scale chemical process. The LiH is remixed with the slurry fluid and transported back to refueling stations scattered over a large area as needed. The basic energy input to the system is provided at the central plant and can be from a variety of energy sources, including fuels like coal, biomass, natural gas, and petroleum oil. All environmental emissions occur at the central processing plant. The vehicle is zero emission, with no hydrocarbon, CO, or CO₂ emissions. The central plant can include more sophisticated emission cleanup processes than would be possible for an onboard processing system.

An important concept feature that needs to be pointed out is the recovery and recycle of the spent hydride at centralized processing plants using a low cost fuel, such as coal or biomass. Regeneration process analysis has indicated that recycling can be performed utilizing a carbothermal process with minimum energy input and at a low cost. Compared to current hydrogen costs of about \$9.00 to \$25.00 per million Btu, this concept should enable hydrogen costs as low as \$3.00 per million Btu to be realized for a LiH system^(1&2). Also, because the hydride reaction will liberate only pure hydrogen, fuel cell catalyst life should be maximized, resulting in high system performance and reliability.

PROTOTYPE DEVELOPMENT

The major objectives of this hydrogen generation development effort are twofold. The first is to use a laboratory-scale system to determine optimum materials and hydrogen generation process conditions to achieve high specific energy for hydrogen supply. The second objective is the design and fabrication of a prototype hydrogen generation system capable of supplying 3.0 kg/hr of high purity hydrogen for fuel cells.

Although there are numerous metal hydrides and organic carrier candidate materials, only a limited number of metal hydrides and organic carrier materials can be used to satisfy DOE's goals of specific weight and volume. One of the essential considerations for the metal hydride is its hydrogen generation efficiency, which includes reaction chemistry between metal hydride and water to complete hydrolysis reactions in a safe and controlled manner. The organic carriers should be chemically inert toward metal hydrides and spent hydrides for storing and transporting, and during hydrolysis reaction. These materials also should be easily separated from spent hydrides, either thermally or mechanically, and be recycled for reuse. Although regeneration of the spent hydrides is not part of the technical effort of this program, it is an important issue for economical and commercial development of the technology.

In the initial effort, we thoroughly analyzed, both theoretically and experimentally, the reaction chemistry of a variety of metal hydrides and water, and the chemical stability of the organic carriers in contact with metal hydrides and spent hydrides. Since detailed hydrolysis reaction kinetics of the metal hydride/organic carrier slurry were not known, we conducted experiments using a high-pressure (2000 psi) and high-temperature (232°C) vessel with temperature, pressure, and magnetic stirrer control capabilities (500 cm³ internal volume). For this investigation, we selected the candidate materials based on the guidelines listed in Table 2.

In the development of a hydride/water activation system, several ideas were considered. These are:

- Single Tank Reactor
- Slurry Atomization Reactor
- Water Bathed Reactor
- Auger-aided Water Vaporizing Reactor

The single tank reactor, shown in Figure 4, is the simplest system. However, several problems exist for this system. The heat exchanger allows hot spots, increasing hydrocarbon contamination. It will also have a slow response to H_2 demand. Furthermore, it is likely that not all hydride will react, leaving a hazardous waste product, and a large volume containing pressurized hydrogen.

The atomized slurry reactor, shown in Figure 5, was conceptualized to remove heat from $15\mu m$ droplets by direct hydrogen heat transfer. This system is complicated and has a wear-prone slurry atomization system. The $\frac{1}{2} m^3/sec$ H_2 flow rates needed for cooling are quite high. In addition, the heat exchanger may be fouled by dust. There is also the likelihood of poor hydride/water mixing reducing generation efficiency, and a large pressurized hydrogen volume is required.

In the water-bathed reactor configuration, shown in Figure 6, heat is removed by the recirculated flow of water. Excess water assures low reaction temperature and complete reaction in a relatively small water to air heat exchanger. Problems, such as the water soaked LiOH waste product and the weight of wasted water, push this concept outside the system goals. The water could be separated by a filter or a filter press, but neither a filter or filter press system allows the concept to reach the weight goals. Also, unfiltered particles will wear the recirculation pump.

In the auger-aided reactor, shown in Figure 7, reactants are pumped to a mixing auger. At 300 rpm, the auger mixes, crushes particles, and eliminates foaming within the hydrogen generation reactor. The waste product contains 10% by mass of water and is a dry, free-flowing powder. About three times the stoichiometric water is added and vaporized by the heat of reaction to control the temperature.

The hydrogen water vapor content in the auger-aided reactor depends on the heat exchanger outlet temperature. Vapor condensation is slowed by the presence of hydrogen, increasing the size of the heat exchanger. The water vapor content could also be reduced by using hydride as a desiccant. This hydrogen production system device achieves the weight and volume goals.

Based on the preliminary analysis and testing of the various concepts discussed above, a prototype system to produce up to 3 kg/hr of hydrogen was designed. This system is shown in Figure 8. To produce the hydrogen, 0.5 l/min 60% LiH slurry flows into the auger reactor, along with 1.4 l/min water for reaction and vapor cooling. The system produces up to .75 kg hydrogen per run. A 1.6 gallon reservoir of 60% LiH slurry, a 5.5 gallon water reservoir, and a 12 gallon hydroxide container make up the reactant and product volumes. A computer controls the hydride and water pumps. Data acquisition of pertinent pressures, temperatures, hydrogen flow, hydrocarbon, and water vapor content are recorded. The system is self-contained on a rolling cart.

A valveless ceramic piston pump is used for the LiH slurry and a gear pump is used for water flow. Three heat exchanger cores with 8-10" fans are used to condense the water from the hydrogen. Table 3 summarizes the energy, mass, and volumetric densities for the system, assuming storage of 15 kg H_2 . To meet the design goals of 3355 Whr/kg and 929 Whr/l, our system must weigh less than 179.9 kg and must have a volume of less than 649.6 l. Table 3 shows that the system designed will meet the weight goal and exceed the volume goal.

SUMMARY

In summary, the following can be stated:

- A hydride/water activation process (the hydrogen generation reactor) has been developed.
- Thermal management design for prototype system has been established.
- A chemical hydride slurry can be used to generate hydrogen for transportation vehicle applications.
- The system has the potential to be safe and easy to use.
- Chemical hydride-based systems can achieve DOE's energy density goals.

REFERENCES

1. Breault, Ronald, "Advanced Chemical Hydride Hydrogen Generation/Storage System for PEM uel Cell Vehicles," to be published in the DOE Office of Advanced Automotive Technology FY 1999 Annual Progress Report.

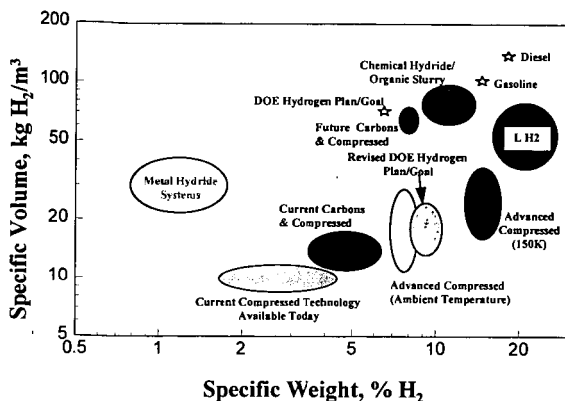


FIGURE 1. Summary of Current and Future Hydrogen Storage Systems

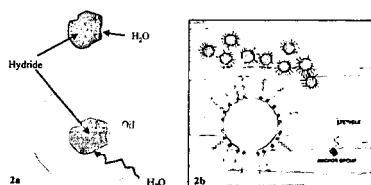


FIGURE 2. Hydride-Water Reaction Concept

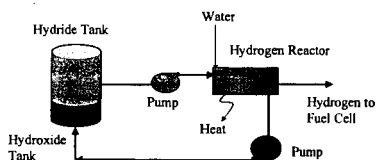


FIGURE 3. Hydrogen - Hydride Concept

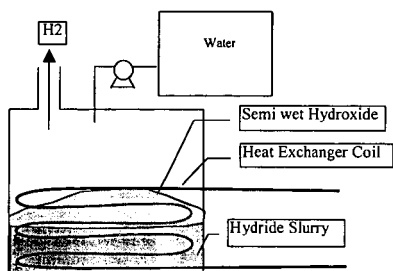


FIGURE 4. Simple One Tank Concept

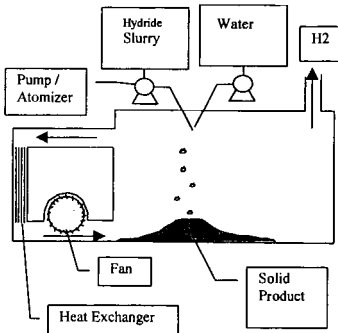


FIGURE 5. Atomized Slurry Reactor Concept

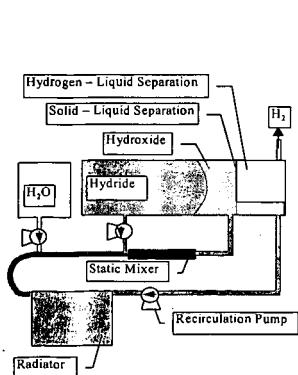


FIGURE 6. Water Bathed Reactor Concept

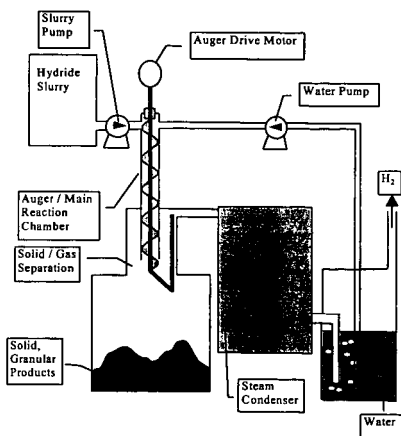


FIGURE 7. Auger-Aided Reactor Concept



| Storage Technology | Specific Weight (HHV) | | Specific Volume (HHV) | | Remarks |
|---|-----------------------|-----------------|-----------------------|-----------------------------------|--|
| | Wh/kg | %H ₂ | Wh/L | kg H ₂ /m ³ | |
| DOE Goal | | | | | -DE-RA02-97EE50443 |
| • Liquid/Gas | 3963/5323 | 9.9/13.4 | 1100/828 | 28/21 | |
| Liquid H₂ | | | | | -Not including boil-off loss |
| • Cryogenic | 6350 | 16.1 | 1250 | 32 | |
| Gaseous H₂ | | | | | -Could be better with new high-pressure tanks |
| • 5000 psia | 2630 | 6.7 | 780 | 20 | -New materials with better capacities |
| Carbon Adsorption | | | | | |
| • 794 psi at 78°K | 2858 | 7.2 | 1535 | 39 | -Need more fundamental research |
| Liquid Hydride | | | | | |
| • Methylcyclohexane | 2070 | 5.9 | 1618 | 46 | |
| Proposed Chemical Hydride Slurry | | | | | -Includes weight and volume of the container, and ancillary components |
| • CaH ₂ | 2670 | 6.8 | 2430 | 62 | |
| • LiH | 5050 | 12.8 | 2430 | 62 | -Does not include reactant water, which is assumed to be provided partially from exhaust gas |
| • NaBH ₄ | 4760 | 12.1 | 2570 | 65 | |
| • LiBH ₄ | 6350 | 16.1 | 2640 | 67 | |

| Metal Hydrides | Organic Carriers |
|--|--|
| <ul style="list-style-type: none"> High specific energy density High hydrogen generation efficiency Relatively inert during storage before and after reaction with water Ease of regeneration Low costs | <ul style="list-style-type: none"> Non-reactive with metal hydrides and spent hydrides Low molecular weight Easy to recycle (easy to separate from spent hydride and water, and to collect for reuse) |

| | | Wt (kg) | Vol (L) |
|---|------------------------------------|---------|---------|
| 65% Lithium Hydride Slurry | | 95.5 | |
| Hydroxide | Direct Gas to Air HX Copper Tube, | | |
| Heat Exchanger | Aluminum Fin | 31.38 | 82.5 |
| Hydride Tank | Stainless | 20 | 120 |
| Hydroxide Tank | Plastic | 12 | 310 |
| Hydride Metering Pump | Valveless Piston | 6.8 | 4 |
| Water Metering Pump | Gear Pump + DC Control | 3 | 2.5 |
| Auger Drive Motor | 1/8 Hp, 5.5:1 Bodine DC Gear Motor | 4 | 2.3 |
| Auger Construction | SS Materials | 5 | 10 |
| Total | | 177.7 | 531.3 |
| System Goals (kg, l) | | 179.9 | 649.6 |
| System Values (watt-hr/kg, watt-hr/liter) | | 3397 | 1136 |
| Goals (watt-hr/kg, watt-hr/liter) | | 3355 | 929 |